Application No. 10/643,751

Amendment Dated August 7, 2006 Reply to Office Action of June 7, 2006

Attorney Docket No. 0116-031357

AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in

the application:

Listing of Claims

Claim 1 (Currently Amended): An electrospray mass spectrometer fitted with

an electrospray ion source, said ion source being provided with comprising a structure

supporting a nebulization nozzle, a sampling orifice, a heated desolvation chamber and a

control knob supporting a support rod, said nebulizing nozzle having an axis and having a

said sampling orifice having an axis, the axis of the nebulization nozzle intersecting the axis

of the sampling orifice, said electrospray ion source further comprising[[:]] a relatively

movable eold-spray desolvation chamber having a direction-changing channel and being

supported from said support rod such that the movable desolvation chamber can be moved off

the axis of the nebulization nozzle in an electrospray ionization mode and set on the axis of

the nebulization nozzle in a cold-spray ionization mode wherein liquid droplets are

introduced from an opening from the nebulization nozzle and pass through the direction-

changing channel such that sample ions are discharged from an exit opposite to the sampling

orifice.

Claim 2 (Original): The electrospray mass spectrometer of claim 1, wherein

said nebulization nozzle consists of a capillary for guiding a sample solution supplied from a

sample inlet port and a pipe for guiding a nebulizing gas introduced from a gas inlet port, said

pipe coaxially surrounding the outer surface of said capillary.

Claim 3 (Currently Amended): The electrospray mass spectrometer of claim 2,

wherein the temperature of said nebulizing gas is adjustable to-room temperature in the

electrospray ionization mode and between room temperature and approximately -50°C in for

use of the electrospray mass spectrometer in the cold-spray ionization mode.

Claim 4 (Currently Amended): The electrospray mass spectrometer of claim 1,

wherein said heated desolvation chamber is cylindrical and said nebulization nozzle is

substantially coaxially inserted in a heated cylindrical desolvation chamber, the nozzle

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opening into the cylindrical desolvation chamber, and wherein said cylindrical desolvation

chamber has a gas inlet port for introducing a heating-and-drying gas.

Claim 5 (Original): The electrospray mass spectrometer of claim 4, wherein a

potential difference of 1-3 kV is imposed between said nebulization nozzle and the sampling

orifice, and wherein a potential difference from zero to hundreds of volts is imposed between

said cylindrical desolvation chamber and the sampling orifice.

Claim 6 (Cancelled).

Claim 7 (Original): The electrospray mass spectrometer of claim 1, wherein

when a mixture of droplets of a sample and a nebulizing gas are electrostatically sprayed

from said nebulization nozzle and the flow rate of sample solution is setable to 1-1,000

microliters per minute.

Claim 8 (Original): The electrospray mass spectrometer of claim 4, wherein in

the electrospray ionization mode, the heating-and-drying gas is introduced into said

cylindrical desolvation chamber from the gas inlet port, and wherein the introduced heating-

and-drying gas and heating performed by a heater buried in an inner wall of the desolvation

chamber cooperate to dry and desolvate the liquid droplets.

Claim 9 (Currently Amended): The electrospray mass spectrometer of claim

8, wherein the heater for the cylindrical desolvation chamber is controllable to approximately

+100 to 300°C between 100 and 300°C.

Claim 10 (Currently Amended): The electrospray mass spectrometer of claim

8, wherein the temperature of said heating-and-drying gas is controllable to approximately

+100 to 300°C between 100 and 300°C.

Claim 11 (Currently Amended): The electrospray mass spectrometer of claim

4, wherein in the cold-spray ionization mode, means for cutting off the supply of the heating-

and-drying gas from the gas inlet port may be cut off and means to deenergize the heater

buried in the inner wall of the cylindrical desolvation chamber deenergized are provided to

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avoid heating of the liquid droplets passing therethrough in the cold-spray mode to thereby

produce multiply charged molecular ions with solvent molecules attached.

Claim 12 (Currently Amended): The electrospray mass spectrometer of claim

11, wherein in the cold-spray ionization mode, means are provided for supplying a cooled gas

may be supplied into said cylindrical desolvation chamber from the gas inlet port.

Claim 13 (Original): The electrospray mass spectrometer of claim 11 or 12,

wherein temperature of said movable desolvation chamber is setable to room temperature or

below in the cold-spray ionization mode.

Claim 14 (Currently Amended): The electrospray mass spectrometer of claim

1, wherein further comprising means for setting the temperature of said desolvation chamber

is setable between room temperature and approximately 0°C in the cold-spray ionization

mode.

Claim 15 (Original): The electrospray mass spectrometer of claim 1, wherein

said cold spray desolvation chamber has a direction-changing channel, and wherein liquid

droplets are introduced from an opening at a side of the nebulization nozzle and passed

through the channel such that sample ions are discharged from an exit opposite to the

sampling orifice.

Claim 16 (Original): The electrospray mass spectrometer of claim 1, wherein

said movable desolvation chamber is supported by a thin support rod for heat insulation.

Claim 17 (Original): The electrospray mass spectrometer of claim 1, wherein

said movable desolvation chamber is fitted with temperature control means such as a

microheater, Peltier element, or sensor.

Claim 18 (Original): The electrospray mass spectrometer of claim 1, wherein

a potential difference of zero to hundreds of volts is developed between said movable

desolvation chamber and said sampling orifice.

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Claim 19 (Cancelled).

Claim 20 (Currently Amended): The electrospray mass spectrometer of claim

1, wherein further comprising means for setting the sampling orifice is setable to a

temperature of approximately +80°C in the electrospray ionization mode and to around room

temperature in the cold-spray ionization mode.

Claim 21 (Original): The electrospray mass spectrometer of claim 1, wherein

the ratio of the amount of ions relative to sample concentration produced in the cold-spray

ionization mode is 1/100 to 1/1,000 of the amount of ions relative to sample concentration

produced in the electrospray ionization mode.

Claim 22 (New): The electrospray mass spectrometer of claim 1, wherein the

direction-changing channel in the movable desolvation chamber is configured to pulverize

the liquid droplets minutely to cause partial vaporization without heating the droplets.

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